## IN THE CLAIMS

- 1. (original): A method of increasing the effective intracellular concentration of a therapeutic molecule within a cell expressing a  $P2X_7$  receptor, the method comprising the step of contacting the cell with the therapeutic molecule and with a substance which modulates the  $P2X_7$  receptor.
- 2. (original): A method according to claim 1, wherein the method comprises the step of contacting the cell with the therapeutic molecule and with a substance which stimulates the P2X<sub>7</sub> receptor.
- 3. (original): A method according to claim 1 or 2, wherein the therapeutic molecule is a cytotoxic drug and it is desired to kill the cell.
- 4. (currently amended): A method according to any one of the preceding claims claim 1, wherein the P2X<sub>7</sub> receptor modulating substance comprises ATP, an analogue of ATP, or an immunoglobulin or immunoglobulin-like variant which possesses specific binding activity for the P2X<sub>7</sub> receptor.
- 5. (currently amended): A method according to any one of the preceding claims, claim 1 wherein the cell expresses an efflux protein which is inhibited by stimulation of the P2X<sub>7</sub> receptor.
- 6. (original): A method according to claim 5, wherein the efflux protein which is inhibited is one or more selected from the group consisting of: P-glycoprotein; mitoxantrone resistance protein; and a member of the multidrug-resistance associated family of proteins.
- 7. (currently amended): A method according to any one of the preceding claims claim 1, wherein the therapeutic molecule and the P2X<sub>7</sub> receptor modulating substance are co-administered.

- 8. (currently amended): A method according to any one of the preceding claims claim 1, wherein the cell is contacted with a substance which inhibits the activity and/or expression of CD45.
- 9. (currently amended): An *in vitro* method in accordance with <del>any of the preceding claims claim 1</del>.
- 10. (canceled)
- 11. (canceled)
- 12. (canceled)
- 13. (canceled)
- 14. (original): A pharmaceutical composition for administration to a mammalian subject, the composition comprising: a therapeutic drug; a P2X<sub>7</sub> receptor modulating substance; and a physiologically acceptable carrier, diluent or excipient.
- 15. (original): A pharmaceutical composition in accordance with claim 14, the composition comprising: a therapeutic drug; a P2X<sub>7</sub> receptor stimulating substance; and a physiologically acceptable carrier, diluent or excipient.
- 16. (original): A pharmaceutical composition according to claim 14 or 15, further comprising a substance which inhibits the activity and/or expression of CD45.
- 17. (original): A method of making a pharmaceutical composition comprising the step of combining in a mixture a therapeutic drug, a P2X<sub>7</sub> receptor modulating substance, and a physiologically acceptable carrier, diluent or excipient.
- 18. (original): A method of making a pharmaceutical composition in accordance with claim 17, the method comprising the step of combining in a mixture a therapeutic drug, a P2X<sub>7</sub> receptor stimulating substance, and a physiologically acceptable carrier diluent or excipient.

- 19. (currently amended): A method according to claim 17 or 18, comprising further combining the ingredients recited in claim 17 or 18 with including a substance which inhibits the activity and/or expression of CD45.
- 20. (original): A method of inhibiting the action of a cell membrane efflux protein, the method comprising the step of contacting a cell expressing a  $P2X_7$  receptor with a substance which causes activation of the  $P2X_7$  receptor.
- 21. (original): An *in vitro* method of inhibiting the action of a cell membrane efflux protein in accordance with claim 20.
- 22. (new): A method of rearranging at least part of the lipid, phospholipid or glycolipid component of a cell membrane which comprises contacting said cell membrane with a  $P2X_7$  receptor modulating substance.